

NOZIN NASAL SANITIZER- alcohol liquid

Denison Pharmaceuticals, LLC

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Nozin Nasal Sanitizer

Active ingredient

Alcohol 62%

Inactive ingredients

Jojoba, water, orange oil, lauric acid, benzalkonium chloride, vitamin E

Purpose

Antiseptic

Use

- to decrease bacteria on the skin

Warnings

For external use only

Flammable. Keep away from fire or flame.

Do not use

- as nose spray
- in eyes
- on mucous membranes
- if you have a history of nasal bleeding or irritation
- if you have allergies to any of the ingredients
- more than four times a day

Stop use and ask a doctor if irritation and redness develop and persist for more than 72 hours.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

adults and children over 12 years of age

- shake bottle very well and remove cap
- apply 2 to 4 drops of solution to the cotton swab
- gently apply wetted cotton to skin inside the rim of one nostril. Swab around nostril rim 6x in each direction. Reapply 2 to 4 drops of solution to cotton swab. Swab around other nostril rim 6x in each direction. Caution: Do not extend the swab into nose beyond swab tip (about 1 cm or 3/8"). Apply to skin only. Swab stem should never enter the nose. Discard swab. Secure cap on bottle.

children under 12 years of age

should be supervised in use

children under 2 years of age

ask a doctor

Other information

- store in a cool, dry place between 15°C - 29°C (59°F - 84°F)



Cyan - Magenta - Yellow - Black - No Coat

Virginia Bessonnier

Director of Operations 2019-08-13

NOZIN NASAL SANITIZER

alcohol liquid

Product Information

| | | | |
|-------------------------|----------------|--------------------|---------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:0295-9025 |
| Route of Administration | TOPICAL | | |

Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|--|-------------------|-----------------|
| ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M) | ALCOHOL | 62 mL in 100 mL |

Inactive Ingredients

| Ingredient Name | | | Strength | |
|---|------------------|--|----------------------|--------------------|
| ORANGE OIL (UNII: AKN3KSD11B) | | | | |
| LAURIC ACID (UNII: 1160N9NU9U) | | | | |
| BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) | | | | |
| .ALPHA.-TOCOPHEROL (UNII: H4N855PNZ1) | | | | |
| WATER (UNII: 059QF0KO0R) | | | | |
| SIMMONDSIA CHINENSIS WHOLE (UNII: DFM16KFA82) | | | | |
| | | | | |
| Packaging | | | | |
| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
| 1 | NDC:0295-9025-62 | 12 mL in 1 BOTTLE; Type 0: Not a Combination Product | 04/09/2020 | |
| | | | | |
| Marketing Information | | | | |
| Marketing Category | | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
| OTC monograph not final | | part333A | 04/09/2020 | |

Labeler - Denison Pharmaceuticals, LLC (001207208)

| Establishment | | | |
|------------------------------|---------|-----------|------------------------|
| Name | Address | ID/FEI | Business Operations |
| Denison Pharmaceuticals, LLC | | 001207208 | manufacture(0295-9025) |

Revised: 4/2020

Denison Pharmaceuticals, LLC